From the INTERNATIONAL SEARCHING AUTHORITY	PERATION TREATY	
To: LEE, Won-Hee 8th Fl.Sung-ji Heights, II 642-16 Yoksam-dong Kangnam-ku Seoul 135-080 Republic of Korea	PCT  WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY  (PCT Rule 43bis.1)	
	Date of mailing (day/month/year) 30 MARCH 2005 (30.03.2005)	
Applicant's or agent's file reference 4FPO-11-04	FOR FURTHER ACTION	
International application No.  PCT/KR2004/003435  International filing da 24 DECEMBER		
International Patent Classification (IPC) or both national classifi	R 2004 (24.12.2004) 27 DECEMBER 2003 (27.12.2003) Cication and IPC.	
IPC7 C07D 307/68		
Applicant		
KOREA RESEARCH INSTITUTE OF CHEMIC	TAI. TECHNOLOGY at al	
. This opinion contains indications relating to the following ite		
Box No. I Basis of the opinion	ems:	
Box No. II Priority		
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Box No. IV Lack of unity of invention	ard to novelty, inventive step and industrial applicability	
Box No. V Reasoned statement under Rule 43bis. 10	Reasoned statement under Rule 43bis 1(a)(i) with regard to povelty, invention etc.	
citations and explanations supporting suc	ch statement	
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Box No. VII Certain defects in the international appl		
Box No. VIII Certain observations on the international	application	
opinions of this International Searching Authority will not be so If this opinion is, as provided above, considered to be a written of	or that this does not apply where the applicant chooses an Authority tified the International Bureau under Rule 66.1bis(b) that written to considered.  opinion of the IPEA, the applicant is invited to submit to the	
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e and mailing address of the ISA/KR		
Korean Intellectual Property Office 920 Dunsan-dong, Seo-gu, Daejeon 302-701, Republic of Korea	Yoon, Kyung Ae	
mile No. 82-42-472-7140	( RRS & Uh )	

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Telephone No. 82-42-481-5605

# WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No.

PCT/KR2004/003435

	is opinion	
l. With regard to the la which it was filed, ur	anguage, this opinion has been established on the basis of the international unless otherwise indicated under this item.	application in the language in
This opinion h	has been established on the basis of a translation from the original language, which is the language of a translation furnished for the purpo	into the following language ses of international search (under
	1 - 7	
claimed invention, thi	nucleotide and/or amino acid sequence disclosed in the international a his opinion has been established on the basis of:	pplication and necessary to the
a. type of material		
a sequence list	listing ted to the sequence listing	
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contained in th	the international application as filed.	
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_	to the purposes of search.	
In addition, in the	case that more than one version or copy of a sequence listing and/or table r	elating thereto has been
and or running,	, are required statements that the information in the subsequent or additional	nomina in ideast, to the
	as filed or does not go beyond the application as filed, as appropriate, were f	urnished.
dditional comments:		
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## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/KR2004/003435

NO

Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement 1. Statement Novelty (N) Claims YES Claims NO 1-10 Inventive step (IS) Claims YES Claims Industrial applicability (IA) 1-10 Claims YES

### 2. Citations and explanations:

Reference is made to the following documents:

Claims

D1 = US 5627193 A (06. 05. 1997)

D2 = US 6630506 B1 (07. 10. 2003)

D3 = WO 03-101450 A1 (11. 12. 2003)

D4 = WO 99-33460 (08.07.1999)

The present invention relates to furancarbonylguanidine derivatives which can be used as a NHE-1 inhibitor, a preparation method thereof and a pharmaceutical composition comprising the same.

D1 discloses quinoline-4-carbonylguanidine derivative and a preparation method therof and a NHE inhibitor containing the same. D2 discloses acyl guanidines which are used as NHE inhibitors. D3 discloses N-((3-oxo 2,3-dihydro-1H-isoindol-1-yl)acetyl)guanidine derivatives as NHE-1 inhibitors for the treatment of infarction and angina pectoris. D4 discloses acyl guanidine sodium/proton exchange inhibitors and method.

#### 1. Novelty

None of the prior art disclose the compound of formula(1) claimed in the present invention and their property. Therefore, the present invention seems to be novel(PCT Article 33(2)).

#### 2. Inventive Step

Although D1-D5 disclose the compounds showing a similar pharmaceutical activity as the compounds of the present invention, neither structural variation nor combination of different structural features of compounds disclosed therein lead to the structural properties as those described in the present invention. Thus the present invention is regarded as being inventive according to PCT Article 33(3).